

K242051 VersaD Delivery CatheterJan 17, 2025
189 days to decisionK242051 · Product code: **QJP** · Neurology
Source: <https://www.510kdatabase.net/k242051/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous, Neurovasculature (QJP)
Date received	Jul 12, 2024
Decision date	Jan 17, 2025
Days to decision	189 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Unity Medical, Inc.
Location	Edina, MN, US
Contact	Matthew Ogle
510(k) history	2 submissions · 2 cleared · 2025-2025

REGULATORY CONSULTANT

Consulting firm	Regulatory Technology Services, LLC
Contact	Prithul Bom

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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